Bone Matters™ Enrollment Fax Form

To sign up your doctor's office to receive reminders about your treatment, ask your doctor to fill out the top portion of this form.

BONE MATTERS™

prolia® (denosumab) injection

Fax Completed Form to: (866) 452-3356

Physician _____

Office Name _____

Office Address _____

City _____ State _____ ZIP _____

Office Fax _____ Phone _____ NPI # _____

Sign Up Today and we will send you important information about your condition, courtesy reminders to help you stay on track with your treatment and more. Please fill out all the fields below.

*Shaded fields are required.

Please complete the following address information:

First Name* _____ Last Name* _____

E-mail Address* _____ Gender* _____

Mailing Address* _____

City* _____ State* _____ ZIP* _____

Date of Birth (mm/dd/yyyy)* _____ / _____ / _____ Date of most recent Prolia® injection* _____ / _____ / _____

We offer reminders via text and phone call. Which would you like to receive?

Call Reminder ☐ Text Reminder ☐ Phone number _____

(A mobile phone number must be provided for text reminders.)

I understand that by checking one of the boxes above and signing below, I consent to Amgen calling and texting me at the phone number(s) I have provided with promotional communications relating to Amgen products and services and/or my condition or treatment. Amgen may use automatic dialing machines or artificial or prerecorded messages to contact me and may leave a voicemail or SMS/text message (standard text messaging rates may apply). I understand that I am not required to provide this consent as a condition of purchasing any goods or services. Reply STOP to cancel SMS messages.

By signing, I acknowledge I have read and understood Amgen's Privacy Notice and Patient Authorization, that I am legally authorized to consent and that I am providing my consent as the patient or the patient's legal guardian.*

_____ Date* _____

(*We cannot send information without signature and date.)

This program is intended for the limited purpose of delivering courtesy reminder notifications to remind you to schedule your next injection of Prolia®. However, it is up to you and/or your caregiver to ensure you obtain your next injection in a timely manner and in accordance with your doctor's instruction. This program is not intended to be a source of medical advice or treatment and does not replace in any way independent medical advice regarding your diagnosis or treatment. Any questions about medications or health care should be presented to your doctor.

By participating, you agree that you are solely responsible for determining whether the use of this program is right for you, and you agree to release Amgen from any liability relating to your use of this program.

Authorization

I authorize Amgen and its contractors and business partners ("Amgen") to use and/or disclose my personal information, *including my personal health information*, only for the following purposes:

- To operate, administer, enroll me in, and/or continue my participation in the Bone Matters™ Patient Support Program and related activities (welcome kit, postcards, tips to manage your condition);
- **To provide me with informational and marketing materials relating to Prolia® products and services, and/or my condition or treatment; and/or**
- To improve, develop, and evaluate products, services, materials and programs related to my condition or treatment.

I understand that the operation and administration of certain of these services and/or programs may require that Amgen contact me by mail, e-mail, telephone or SMS/text. I understand and consent to Amgen contacting me using the contact information provided in this form to enroll me in, operate, and administer Amgen patient support services and/or programs as described above other than promotional and injection reminder communications by telephone or SMS/text (which I can separately opt-in on the right).

I further understand that the Bone Matters™ Patient Support Program and additional informational and marketing communications related to my condition and treatment are optional and free services. I do not have to sign this authorization and this authorization in no way affects my right to obtain any medications. To obtain a copy of this authorization or opt-out at any time, I can contact Amgen by calling 800-917-1622 or by writing to PO Box 781046, Indianapolis, IN 46278. The Amgen Privacy Statement can be found at www.Prolia.com.

By signing this form above, I agree to enroll in the Bone Matters™ Patient Support Program, and to receive informational and marketing communications from Amgen. If you do not want your information used for the purposes described above, you can opt-out anytime.

If you are under the age of 18, you are not eligible to participate, and we ask you not to submit any personal information to us.

Please see Indications and Important Safety Information on the next page.

Indications

Prolia® is a prescription medicine used to treat osteoporosis in women after menopause who are at high risk for fracture or cannot use another osteoporosis medicine or other osteoporosis medicines did not work well.

Prolia® is a prescription medicine used to increase bone mass in men with osteoporosis who are at high risk for fracture.

Important Safety Information

Do not take Prolia® if you: have low blood calcium; or are pregnant or plan to become pregnant, as Prolia® may harm your unborn baby; or are allergic to denosumab or any ingredients in Prolia®.

What is the most important information I should know about Prolia®?

If you receive Prolia®, you should not receive XGEVA®. Prolia® contains the same medicine as XGEVA® (denosumab).

Prolia® can cause serious side effects:

Serious allergic reactions have happened in people who take Prolia®. Call your doctor or go to your nearest emergency room right away if you have any symptoms of a serious allergic reaction, including low blood pressure (hypotension); trouble breathing; throat tightness; swelling of your face, lips, or tongue; rash; itching; or hives.

Low blood calcium (hypocalcemia). Prolia® may lower the calcium levels in your blood. If you have low blood calcium, it may get worse during treatment. Your low blood calcium must be treated before you receive Prolia®.

Take calcium and vitamin D as your doctor tells you to help prevent low blood calcium.

Severe jaw bone problems (osteonecrosis) may occur. Your doctor should examine your mouth before you start Prolia® and may tell you to see your dentist. It is important for you to practice good mouth care during treatment with Prolia®.

Unusual thigh bone fractures. Some people have developed unusual fractures in their thigh bone. Symptoms of a fracture include new or unusual pain in your hip, groin, or thigh.

Increased risk of broken bones, including broken bones in the spine, after stopping Prolia®. After your treatment with Prolia® is stopped, your risk for breaking bones, including bones in your spine, is increased. Your risk for having more than 1 broken bone in your spine is increased if you have already had a broken bone in your spine. Do not stop taking Prolia® without first talking with your doctor. If your Prolia® treatment is stopped, talk to your doctor about other medicine that you can take.

Serious infections in your skin, lower stomach area (abdomen), bladder, or ear may happen. Inflammation of the inner lining of the heart (endocarditis) due to an infection may also happen more often in people who take Prolia®. You may need to go to the hospital for treatment.

Prolia® is a medicine that may affect the ability of your body to fight infections. People who have weakened immune systems or take medicines that affect the immune system may have an increased risk for developing serious infections.

Skin problems such as inflammation of your skin (dermatitis), rash, and eczema have been reported.

Bone, joint, or muscle pain. Some people who take Prolia® develop severe bone, joint, or muscle pain.

Before taking Prolia®, tell your doctor about all of your medical conditions, including if you:

- Take the medicine XGEVA® (denosumab)
- Have low blood calcium
- Cannot take daily calcium and vitamin D
- Had parathyroid or thyroid surgery (glands located in your neck)
- Have been told you have trouble absorbing minerals in your stomach or intestines (malabsorption syndrome)
- Have kidney problems or are on kidney dialysis
- Plan to have dental surgery or teeth removed
- Are pregnant or plan to become pregnant

  Females who are able to become pregnant:

  - Your healthcare provider should do a pregnancy test before you start treatment with Prolia®.
  - You should use an effective method of birth control (contraception) during treatment with Prolia® and for at least 5 months after your last dose of Prolia®.
  - Tell your doctor right away if you become pregnant while taking Prolia®.
  - Are breast-feeding or plan to breast-feed

What are the possible side effects of Prolia®?

It is not known if the use of Prolia® over a long period of time may cause slow healing of broken bones. The most common side effects of Prolia® in women being treated for osteoporosis after menopause are back pain, pain in your arms and legs, high cholesterol, muscle pain, and bladder infection. The most common side effects of Prolia® in men with osteoporosis are back pain, joint pain, and common cold (runny nose or sore throat).

These are not all the possible side effects of Prolia®. Call your doctor for medical advice about side effects.

You are encouraged to report negative side effects of prescription drugs to the FDA. Visit www.fda.gov/medwatch, or call 1-800-FDA-1088.

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Please see accompanying PROLIA® full Prescribing Information, including Medication Guide.

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